CLAIMS

What is claimed is:

- A method for preventing or significantly reducing a risk of cardiovascular disease in a healthy subject comprising administering an effective dose of an ACE inhibitor to the healthy subject, whereby the risk of cardiovascular disease in the healthy subject is prevented or significantly reduced.
- The method of claim 1, wherein the ACE inhibitor comprises ramipril.
- The method of claim 1, wherein the healthy subject comprises
 a subject free of hypertension, congestive heart failure, left ventricular
 dysfunction, prior myocardial infarct, and induced activation of the reninangiotensin system.
- The method of claim 1, wherein the healthy subject comprises
 a post-menopausal female human subject.
 - 5. The method of claim 1, wherein the healthy subject comprises a subject comprising a PAI-1 polymorphism, wherein the PAI-1 polymorphism results in an elevated level of PAI-1 when compared to a control level of PAI-1.
- The method of claim 5, wherein the PAI-1 polymorphism comprises a 4G PAI-1 polymorphism.
 - 7. The method of claim 1, wherein reducing the risk of cardiovascular disease in the healthy subject comprises significantly reducing a plasma level of PAI-1 in the healthy subject.

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- The method of claim 1, wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 35% compared to a baseline plasma level of PAI-1.
- 9. The method of claim 8, wherein the healthy subject comprises
 5 a subject comprising a PAI-1 polymorphism, wherein the PAI-1 polymorphism is correlated with elevated levels of PAI-1 activity, and wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 35% compared to a baseline plasma level of PAI-1.
 - 10. A method for reducing a plasma level of PAI-1 in a healthy subject comprising administering an effective dose of an ACE inhibitor to the subject, whereby the plasma level of PAI-1 in the subject is reduced.
 - The method of claim 10, wherein the ACE inhibitor comprises ramipril.
- 15 12. The method of claim 10, wherein the healthy subject comprises a subject free of hypertension, congestive heart failure, left ventricular dysfunction, and prior myocardial infarct, and induced activation of the reninangiotensin system.
- The method of claim 10, wherein the healthy subject comprises
 a post-menopausal female human subject.
 - 14. The method of claim 10, wherein the healthy subject comprises a subject comprising a PAI-1 polymorphism, wherein the PAI-1 polymorphism is correlated with an elevated level of PAI-1 when compared to a control level of PAI-1.

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- The method of claim 14, wherein the PAI-1 polymorphism comprises a 4G PAI-1 polymorphism.
- 16. The method of claim 10, wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 35% compared to a baseline plasma level of PAI-1.
- 17. The method of claim 16, wherein the healthy subject comprises a subject comprising a PAI-1 polymorphism, wherein the PAI-1 polymorphism is correlated with elevated levels of PAI-1 activity, and wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 35% compared to a baseline plasma level of PAI-1.
- 18. A method for preventing or significantly reducing a risk of cardiovascular disease in a healthy subject comprising co-administering an effective dose of an ACE inhibitor and an effective dose of hormone, or a conjugate thereof, to the healthy subject, whereby the risk of cardiovascular disease in the healthy subject is prevented or significantly reduced.
- The method of claim 18, wherein the ACE inhibitor comprises ramipril.
- The method of claim 19, wherein the hormone comprises
 PREMARIN® estrogen.
 - 21. The method of claim 18, wherein the healthy subject comprises a subject free of hypertension, congestive heart failure, left ventricular dysfunction, prior myocardial infarct, and induced activation of the reninangiotensin system.

- The method of claim 18, wherein the healthy subject comprises a post-menopausal female human subject.
- 23. The method of claim 18, wherein the healthy subject comprises
 a subject comprising a PAI-1 polymorphism, wherein the PAI-1
 polymorphism is correlated with an elevated level of PAI-1 when compared
 to a control level of PAI-1.
- The method of claim 23, wherein the PAI-1 polymorphism comprises a 4G PAI-1 polymorphism.
- 25. The method of claim 18, wherein reducing the risk of cardiovascular disease in the healthy subject comprises significantly reducing a plasma level of PAI-1 in the healthy subject.
 - 26. The method of claim 25, wherein the significantly reducing a plasma level of PAI-1 in the healthy subject comprises reducing the plasma level of PAI-1, wherein the reducing is to a greater extent when compared to a reducing a plasma of PAI-1 following administration of an ACE inhibitor or hormone to the subject.
 - 27. The method of claim 25, wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 50% when compared to a baseline plasma level of PAI-1.
- 28. A method for reducing a plasma level of PAI-1 in a healthy subject comprising co-administering an effective dose of an ACE inhibitor and an effective dose of hormone to the subject, whereby the plasma level of PAI-1 in the subject is reduced.

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- The method of claim 28, wherein the ACE inhibitor comprises ramipril.
- The method of claim 28, wherein the hormone comprises PREMARIN® estrogen.
- 31. The method of claim 28, wherein the healthy subject comprises a subject free of hypertension, congestive heart failure, left ventricular dysfunction, prior myocardial infarct, and induced activation of the reninangiotensin system.
- The method of claim 28, wherein the healthy subject comprises
 a post-menopausal female human subject.
 - 33. The method of claim 28, wherein the healthy subject comprises
 a subject comprising a PAI-1 polymorphism, wherein the PAI-1
 polymorphism is correlated with an elevated level of PAI-1 when compared
 to a control level of PAI-1
 - The method of claim 28, wherein the PAI-1 polymorphism comprises a 4G PAI-1 polymorphism.
 - 35. The method of claim 28, wherein the significantly reducing a plasma level of PAI-1 in the healthy subject comprises reducing the plasma level of PAI-1, wherein the reducing is to a greater extent when compared to a reducing a plasma of PAI-1 following administration of an ACE inhibitor or estrogen to the subject.
 - 36. The method of claim 28, wherein the significantly reducing a plasma level of PAI-1 comprises reducing a plasma level of PAI-1 by at least about 50% when compared to a baseline plasma level of PAI-1.